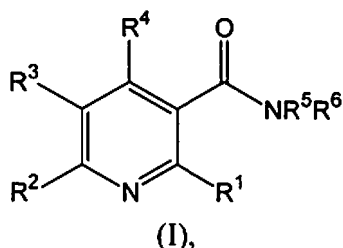


1. (Currently amended) A method of inhibiting angiogenesis **by inhibiting cell migration,**  
**said method** comprising administering to a mammal in need of such treatment a compound of  
 formula (I)



or a therapeutically salt thereof, wherein

$R^1$ ,  $R^2$ ,  $R^3$ , and  $R^4$  are independently selected from the group consisting of hydrogen, alkoxy, alkoxy carbonyl alkyl, alkyl, amino, aryl, aryl alkyl, cyano, cyano alkyl, cyclo alkyl, (cyclo alkyl) alkyl, halo, halo alkoxy, halo alkyl, hetero aryl, hetero aryl alkyl, heterocycle, (heterocycle) alkyl, hydroxy, hydroxy alkyl, and nitro alkyl; and

$R^5$  and  $R^6$  are independently selected from the group consisting of hydrogen, alkoxy alkyl, alkyl, alkynyl, alkyl sulfanyl alkyl, amino alkyl, cyano alkyl, cyclo alkyl, (cyclo alkyl) alkyl, halo alkyl, hetero aryl alkyl, and (heterocycle) alkyl.

2. (Withdrawn) The method of claim 1 wherein  $R^4$  is hydrogen.

Claims 3-17 (Canceled)

18. (Withdrawn) The method of claim 2 wherein

$R^1$  and  $R^2$  are other than hydrogen; and

$R^3$  is hydrogen.

19. (Withdrawn) The method of claim 18 wherein one of  $R^5$  and  $R^6$  is alkyl and the other is selected from the group consisting of hydrogen and alkyl.

20. (Original) The method of claim 19 wherein the compound of formula (I) is selected from the group consisting of

- 19 2-chloro-N-ethyl-N-isopropyl-6-methylnicotinamide;
- 21 2-chloro-N,N,6-trimethylnicotinamide;
- 24 N,N-diethyl-2-methyl-6-(trifluoromethyl)nicotinamide;

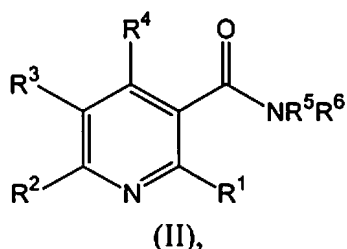
- 103 N-isopropyl-N,2-dimethyl-6-(trifluoromethyl)nicotinamide;
- 104 N-butyl-N,2-dimethyl-6-(trifluoromethyl)nicotinamide;
- 105 N-isobutyl-N,2-dimethyl-6-(trifluoromethyl)nicotinamide;
- 109 N-ethyl-2-methyl-N-propyl-6-(trifluoromethyl)nicotinamide;
- 110 N-butyl-N-isopropyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 112 N-isopropyl-2-methyl-N-propyl-6-(trifluoromethyl)nicotinamide;
- 113 N-butyl-2-methyl-N-propyl-6-(trifluoromethyl)nicotinamide;
- 115 N,N-dibutyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 116 N,N-diisobutyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 119 N-(sec-butyl)-2-chloro-6-methylnicotinamide;
- 120 2-chloro-6-methyl-N-pentylnicotinamide;
- 121 2-chloro-6-methyl-N-(2-methylbutyl)nicotinamide;
- 135 N-ethyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 136 2-methyl-N-propyl-6-(trifluoromethyl)nicotinamide;
- 137 N-isopropyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 138 N-butyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 139 N-(sec-butyl)-2-methyl-6-(trifluoromethyl)nicotinamide;
- 140 N-isobutyl-2-methyl-6-(trifluoromethyl)nicotinamide;
- 141 N-(tert-butyl)-2-methyl-6-(trifluoromethyl)nicotinamide;
- 142 2-methyl-N-pentyl-6-(trifluoromethyl)nicotinamide;
- 143 2-methyl-N-(1-methylbutyl)-6-(trifluoromethyl)nicotinamide;
- 144 2-methyl-N-(2-methylbutyl)-6-(trifluoromethyl)nicotinamide;
- 145 2-methyl-N-(3-methylbutyl)-6-(trifluoromethyl)nicotinamide;
- 146 N-(1,1-dimethylpropyl)-2-methyl-6-(trifluoromethyl)nicotinamide;
- 147 N-(1-ethylpropyl)-2-methyl-6-(trifluoromethyl)nicotinamide;
- 148 N-hexyl-2-methyl-6-(trifluoromethyl)nicotinamide; and
- 149 N-(3,3-dimethylbutyl)-2-methyl-6-(trifluoromethyl)nicotinamide.

Claims 21-24 (Canceled).

25. (Currently amended) A method of inhibiting angiogenesis by inhibiting cell migration, said method comprising administering to a human in need of such treatment a compound of claim 1 or a therapeutically acceptable salt thereof.

Claims 26-30 (Canceled)

31. (Currently amended) A method of inhibiting angiogenesis **by inhibiting cell migration,** **said method** comprising administering to a mammal in need of such treatment a compound of formula (II)



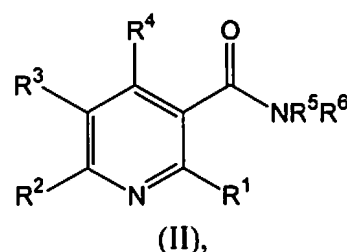
or a therapeutically acceptable salt thereof, wherein

$R^1$  and  $R^4$  are independently selected from the group consisting of hydrogen, alkoxy, alkoxycarbonylalkyl, alkyl, arylalkyl, cyano, cyanoalkyl, cycloalkyl, (cycloalkyl)alkyl, halo, haloalkoxy, haloalkyl, heteroaryl, heteroarylalkyl, heterocycle, (heterocycle)alkyl, hydroxy, hydroxyalkyl, and nitroalkyl;

$R^2$  and  $R^3$  are independently selected from the group consisting of hydrogen, alkoxy, alkoxycarbonylalkyl, alkyl, aryl, arylalkyl, cyano, cyanoalkyl, cycloalkyl, (cycloalkyl)alkyl, halo, haloalkoxy, haloalkyl, heteroaryl, heteroarylalkyl, heterocycle, (heterocycle)alkyl, hydroxy, and hydroxyalkyl; provided that at least one of  $R^1$ ,  $R^2$ ,  $R^3$ , and  $R^4$  is other than hydrogen; and

one of  $R^5$  and  $R^6$  is alkyl and the other is selected from the group consisting of alkoxyalkyl and dialkylaminoalkyl.

32. (Currently amended) A method of inhibiting angiogenesis **by inhibiting cell migration,** **said method** comprising administering to a human in need of such treatment a compound of formula (II)



or a therapeutically acceptable salt thereof, wherein

$R^1$  and  $R^4$  are independently selected from the group consisting of hydrogen, alkoxy, alkoxycarbonylalkyl, alkyl, arylalkyl, cyano, cyanoalkyl, cycloalkyl, (cycloalkyl)alkyl, halo, haloalkoxy, haloalkyl, heteroaryl, heteroarylalkyl, heterocycle, (heterocycle)alkyl, hydroxy, hydroxyalkyl, and nitroalkyl;

$R^2$  and  $R^3$  are independently selected from the group consisting of hydrogen, alkoxy, alkoxycarbonylalkyl, alkyl, aryl, arylalkyl, cyano, cyanoalkyl, cycloalkyl, (cycloalkyl)alkyl,

halo, haloalkoxy, haloalkyl, heteroaryl, heteroarylalkyl, heterocycle, (heterocycle)alkyl, hydroxy, and hydroxyalkyl; provided that at least one of  $R^1$ ,  $R^2$ ,  $R^3$ , and  $R^4$  is other than hydrogen; and one of  $R^5$  and  $R^6$  is alkyl and the other is selected from the group consisting of alkoxyalkyl and dialkylaminoalkyl.

Claims 33-34 (Canceled)